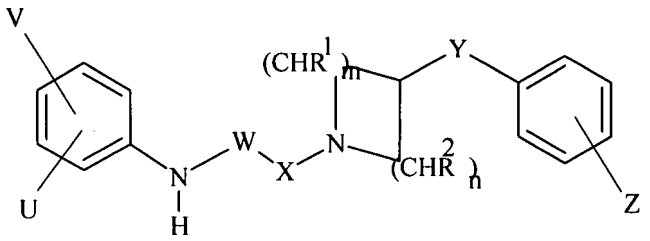
Amendments to the Claims

The listing of claims will replace all prior versions, and listing of claims in the application.

1. (Currently Amended) A compound comprising a structure of formula (I):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof.

2. (Original) The compound of claim 1 wherein $-(CHR^1)_m$ and $-(CHR^2)_n$ are each $-CH_2-CH_2-$.

3. (Original) The compound of claim 2 wherein W is $-CH_2-$.

4. (Original) The compound of claim 2 wherein W is -CO-.

5. (Canceled)

6. (Canceled)

7. (Canceled)

8. (Canceled)

9. (Canceled)

10. (Canceled)

11. (Canceled)

12. (Canceled)

13. (Canceled)

14. (Canceled)

15. (Previously Presented) The compound of claim 1 wherein the compound is:

2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-lH-indol-5-yl)--acetamide;

2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)-acetamide;

2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-lH benzimidazol-5-yl) acetamide;

2-(4-benzyl-piperidin-l-yl)-2-oxo-N-(2-oxo-2,3-dihydro-lH-indol-5-yl)-acetamide;

2-(4-benzyl-piperidin-1-yl)-2-oxo-N-(2-oxo-2,3-dihydro-lH-benzimidazol-5-yl)-acetamide;

2-(4-benzyl-piperidin-1-yl)-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)-acetamide;

5-{2-[4-(4-fluoro-benzyl)-piperidin-l-yl]-2-oxo-ethylamino}-1,3-dihydro-benzoimidazol-2-one;

6-{2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-2-oxo-ethylamino}-3H-benzoxazol-2-one;

2-[4-(4-methylbenzyl)-piperidin-l-yl]-2-oxo-N-(3-oxo-3,4-dihydro-2H-benzo[1,4] oxazin7-yl) acetamide;

2-[4-[4-methyl-benzyl)-piperidin-l-yl]-2-oxo-N-(2-oxo-2,3-dihydro-lH-indol-5-yl)-acetamide;

2-[4-(4-chloro-phenoxy)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-lH-benzimidazol-5-yl) acetamide;

2-[4-(4-chloro-phenoxy)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

2-[4-(4-chloro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)-acetamide;

2-[4-(4-chloro-benzyl)-piperidin-l-yl]-2-oxo-N-(2-oxo-2,3-dihydro-lH-benzimidazol-5yl) acetamide;

2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)-2-(4-p-tolyloxy-piperidin-l-yl)-acetamide;

2-oxo-N-(2-oxo-2,3-dihydro-lH-benzimidazol-6-yl)-2-(4-p-tolyloxy-piperidin-I-yl)-acetamide;

2-[4-(4-chloro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-1 H-indol-5-yl)-acetamide;

6-[2-(4-benzyl-piperidin-1-y1)-2-oxo-ethylamino]-3H-benzoxazol-2-one;

2-(4-benzyl-piperidin-l-yl)-N-(2-mercapto-3H-benzimidazol-5-yl)-2-oxo-acetamide;

2-(4-benzyl-piperidin-1-yl)-2-oxo-N-(2-oxo-2,3-dihydro-benzothiazol-6-yl)-acetamide;

2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-N-(2-mercapto-3H-benzimidazol-5-yl)-2-oxo-acetamide;

2-[4-(4-fluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzothiazol-6-yl) acetamide;

2-oxo-N-(2-oxo-2,3-dihydro-benzothiazol-6-yl)-2-(4-p-tolyloxy-piperidin-l-yl)-acetamide;

N-(2-mercapto-3H-benzimidazol-5-yl)-2-(4-p-tolyloxy-piperidin-l-yl)-2-oxo-acetamide;

2-[4-(4-methyl-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl] acetamide;

2-[4-[4-methoxy-benzyl)-piperidin-1-yi]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

2-[4-[3-methoxy-benzyl)-piperidin-l-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

2-[4-[3-methyl-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

2-[4-(4-cyano-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)acetamide;

2-[4-[3-fluoro-benzyl)-piperidin-1-~~yL~~yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)acetamide;

2-[4-(2,4-difluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

6-(2-[4-(4-methyl-benzyl)-piperidin-l-yl]-2-oxo-ethylamino}-3H-benzoxazol-2-one;

2-[4-(3,4-difluoro-benzyl)-piperidin-1-yl]-2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl) acetamide;

2-[4-(4-methyl-benzyl)-piperidin-1-yl)-2-oxo-N-(2-oxo-1,2,3,4-tetrahydro-quinolin-6-yl acetamide;

2-[4-(4-methyl-benzyl)-piperidin-1-yl)-2-oxo-N-(2-oxo-2,3-dihydro-benzothiazol-6-yl) acetamide;

2-[4-(4-chloro-phenoxy)-piperidin-1-yl]-2-oxo N-(2-oxo-2,3-dihydro-benzothiazol-6-yl) acetamide; or

2-oxo-N-(2-oxo-2,3-dihydro-benzoxazol-6-yl)-2-(4-p-tolyloxy-piperidin-l-yl)-acetamide;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof.

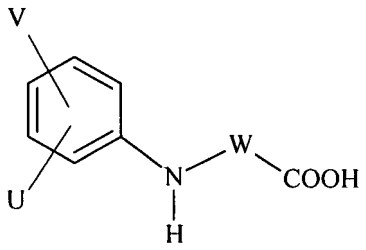
16. (Original) The compound of claim 1 wherein the compound is a functional antagonist of NMDA receptors.

17. (Original) The compound of claim 16 wherein the compound is a functional NR2B subtype specific NMDA receptor antagonist.

18. (Original) The compound of claim 16 wherein the compound exhibits an IC_{50} value of less than 54 μM in a NMDA antagonism or binding test.

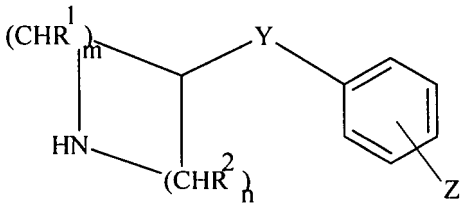
19. (Original) The compound of claim 18 wherein the compound exhibits an IC_{50} value of less than 5 μM in a NMDA antagonism or binding test.

20. (Original) The compound of claim 1, which is synthesized by a method comprising reacting a carboxylic acid of formula (II):



(II)

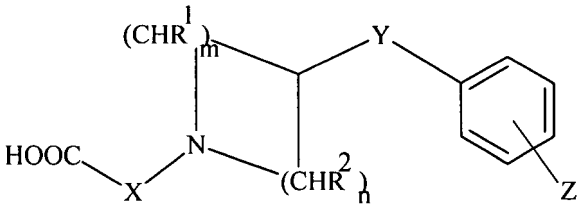
wherein U, V and W are as defined in claim 1, or a reactive derivative thereof, with an amine of formula (III):



(III)

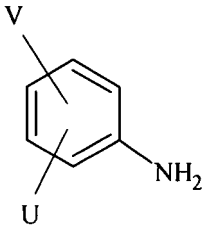
wherein R^1, R^2, Y, Z, n and m are as defined in claim 1.

21. (Original) The compound of claim 1, wherein W is -CO-, synthesized by a method comprising reacting a carboxylic acid of formula (IV):



(IV)

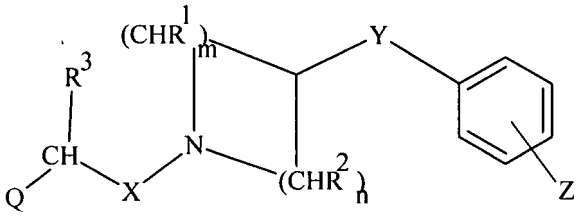
wherein X, R^1, R^2, Y, Z, n and m are as defined in claim 1, or a reactive derivative thereof, with
an amine of formula (V):



(V)

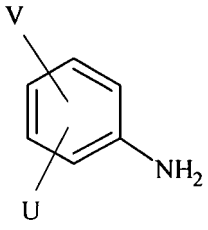
wherein U and V are as defined in claim 1.

22. (Original) The compound of claim 1, wherein W is $-CH_2-$ or $-CH_2-(C_1-C_4$ alkyl)-, synthesized by a method comprising reacting a halogen derivative of formula (VII):



(VII)

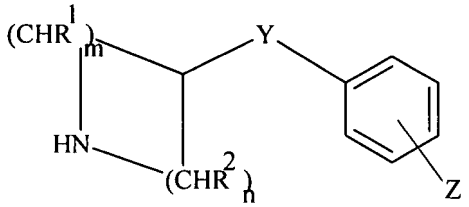
wherein Q is halogen, R^3 is hydrogen atom or a C_1-C_4 alkyl and X, R^1, R^2, Y, Z, n and m are as defined in claim 1 with an amine of formula (V):



(V)

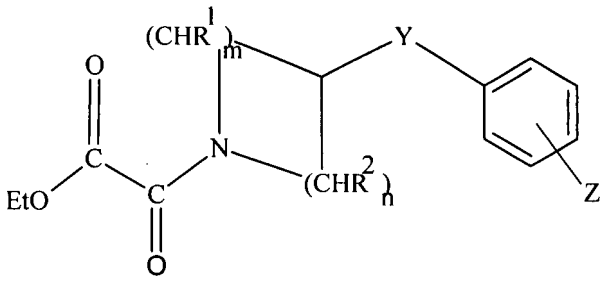
wherein U and V are as defined in claim 1.

23. (Original) The compound of claim 1 synthesized by a method comprising reacting a secondary amine of formula (III):



(III)

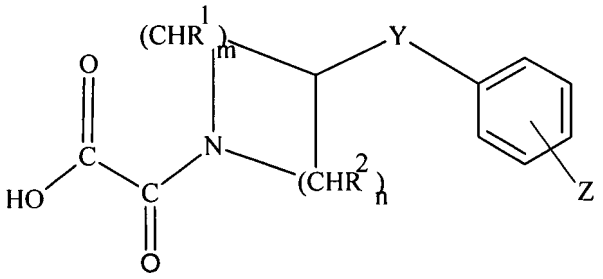
where R^1, R^2, m, n, Y and Z are as defined in claim 1 with ethyl oxalylchloride in the presence of a solid-supported base in dichloromethane to produce an ester compound of formula (VIII):



(VIII)

wherein R', R^2, m, n, Y and Z are as defined as in claim 1;

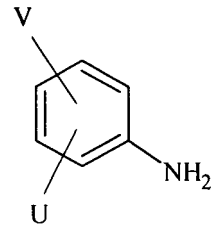
saponifying the ester compound of formula (VIII) with a strongly basic ion exchange resin in ethanol to produce an oxalamid acid of formula (IX):



(IX)

where R^1, R^2, m, n, Y and Z are as defined in claim 1; and

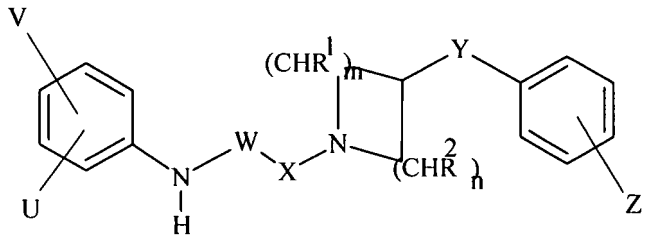
reacting the oxalamid acid of formula (IX) with an amide of formula (V):



(V)

wherein U and V are as defined in claim 1 in a mixture of dichloromethane and dimethylformamide in the presence of 1-[3-(dimethylamino)-propyl]-3-ethylcarbodiimide to produce the compound of claim 1.

24. (Currently Amended) A pharmaceutical composition comprising a biologically effective dose of a compound of formula (1):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, -N(C_1-C_4 alkyl)-, - C_1-C_4 alkylene-N(C_1-C_4 alkyl)-, -CH_2O-, -CH(OH)- or -OCH_2-;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, and one or more pharmaceutical carriers.

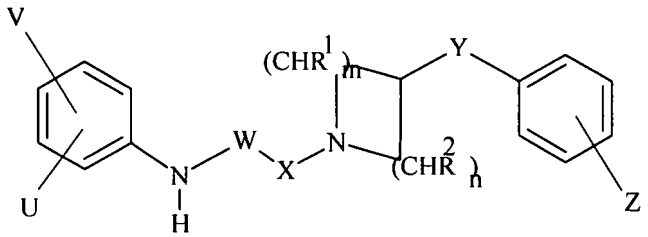
25. (Original) The pharmaceutical composition of claim 24 wherein the compound is a functional antagonist of NMDA receptors.

26. (Original) The pharmaceutical composition of claim 25 wherein the compound is a functional NR2B subtype specific NMDA receptor antagonist.

27. (Original) The pharmaceutical composition of claim 24 wherein the pharmaceutical composition contains 0.01 to 100 mg of the compound in a single dosage unit.

28. (Original) The pharmaceutical composition of claim 24 wherein the pharmaceutical composition is in the form of a tablet.

29. (Currently Amended) A process for synthesizing a compound of formula (I):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

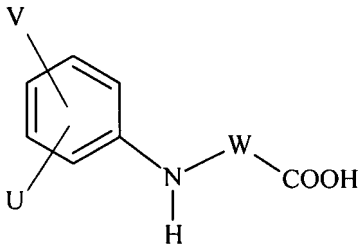
Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

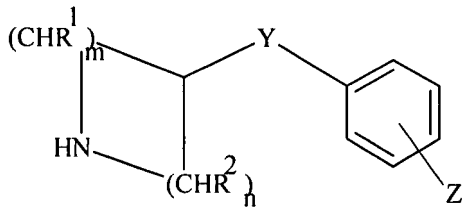
n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, comprising: reacting a carboxylic acid of formula (II):



(II)

wherein U, V and W are as defined above, or a reactive derivative thereof, with an amine of formula (III):

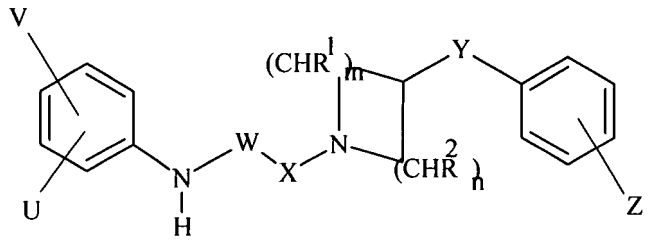


(III)

wherein R^1, R^2, Y, Z, n and m are as defined above.

30. (Original) The process of claim 29 wherein the reactive derivative of the carboxylic acid of formula (II) is formed using O-benzotriazol-1-yl-N,N,N',N' tetramethyluronium hexafluorophosphate.

31. (Currently Amended) A process for synthesizing a compound of formula (1):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is <u>selected from the group consisting of</u> morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

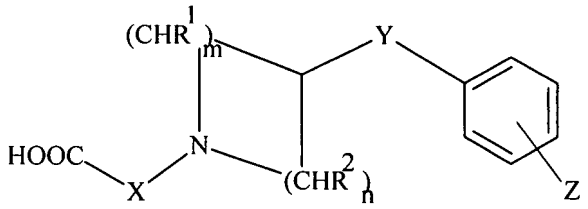
Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

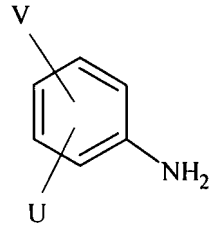
or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, comprising:

reacting a carboxylic acid of formula (IV):



(IV)

wherein X, R^1, R^2, Y, Z, n and m are as defined above, or a reactive derivative thereof, with an amine of formula (V):

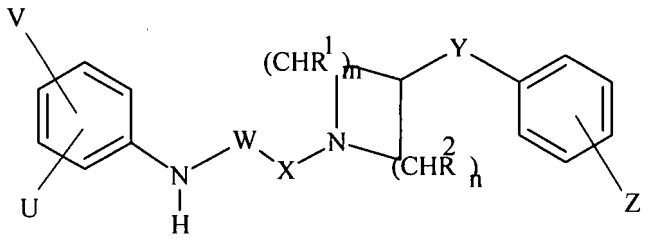


(V)

wherein U and V are as defined above.

32. (Original) The process of claim 31 wherein the reactive derivative of the carboxylic acid of formula (IV) is formed using O-benzotriazol-1-yl-N,N,N,N tetramethyluronium hexafluorophosphate.

33. (Currently Amended) A process for synthesizing a compound of formula (I):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

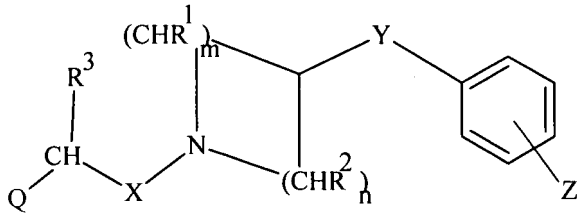
Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

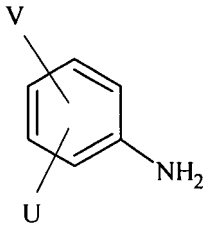
n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, comprising: reacting a halogen derivative of formula (VII):



(VII)

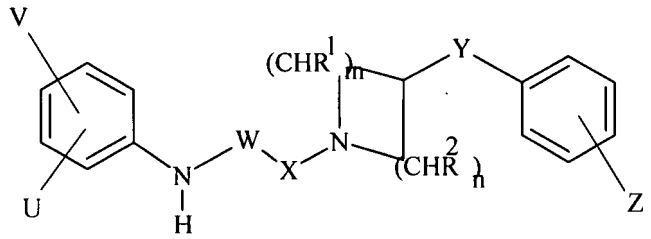
wherein Q is halogen, R^3 is hydrogen or a C_1-C_4 alkyl and X, R^1, R^2, Y, Z, n and m are as defined above with an amine of formula (V):



(V)

wherein U and V are as defined above.

34. (Currently Amended) A process for synthesizing a compound of formula (1):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

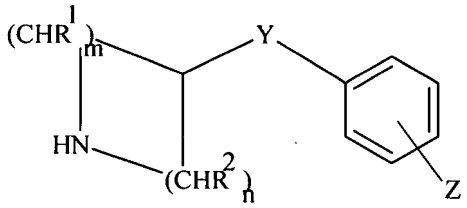
Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

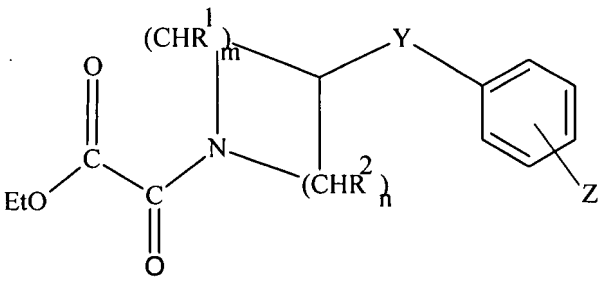
or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, comprising:

reacting a secondary amine of formula (III):



(III)

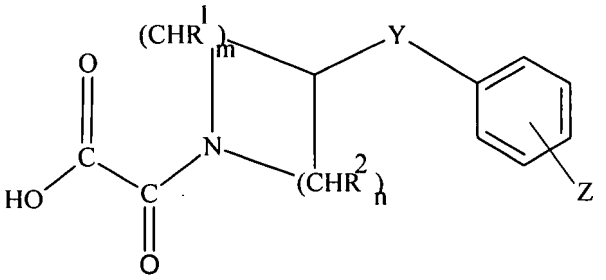
where R', R^2, m, n, Y and Z are as defined above with ethyl oxalylchloride in the presence of a solid-supported base in dichloromethane to produce an ester compound of formula (VIII):



(VIII)

wherein R', R^2, m, n, Y and Z are as defined above,

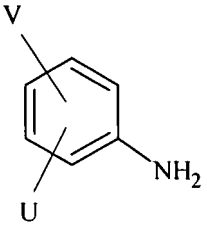
saponifying the ester compound of formula (VIII) with a strongly basic ion exchange resin in ethanol to produce an oxalamid acid of formula (IX):



(IX)

where R', R^2, m, n, Y and Z are as defined above, and

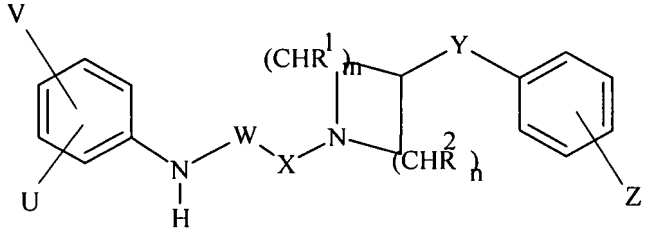
reacting the oxalamid acid of formula (IX) with an amide of formula (V):



(V)

wherein U and V are as defined above in a mixture of dichloromethane and dimethylformamide in the presence of 1-[3-(dimethylamino)-propyl]-3-ethylcarbodiimide to produce the compound of claim 1.

35. (Currently Amended) A process for manufacturing pharmaceutical compositions comprising mixing a compound of formula (1):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

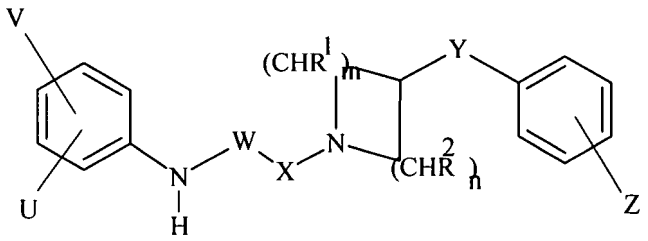
n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof, with a pharmaceutical carrier.

36. (Original) The process of claim 35 wherein the compound is a functional NR2B subtype specific NMDA receptor antagonist.

37. (Canceled)

38. (Withdrawn and Currently Amended) A method for alleviating pain in a mammal comprising administering to the mammal a compound of formula (I):



(I)

wherein:

V and U:

together form a group that contains one or more heteroatoms, and that taken together with one or more:

(a) hydrogen atoms;

(b) carbon atoms;

(c) -CH= groups;

(d) $-CH_2-$ groups; or

(e) additional heteroatoms of the same or different type;

or any combination thereof, form a 4-7 membered homocyclic or heterocyclic ring, wherein the ~~homocyclic or~~ heterocyclic ring is selected from the group consisting of morpholine, pyrrole, pyrrolidine, oxo-pyrrolidine, thioxo-pyrrolidine, pyrazole, pyrazolidine, imidazole, oxo-imidazole, thioxo-imidazole, imidazolidine, oxo-imidazolidine, thioxo-imidazolidine, 1,4-oxazine, oxazole, oxazolidine, oxo-oxazolidine, thioxo-oxazolidine or 3-oxo-1,4-oxazine;

W: is -CO-, $-CH_2-$ or $-CH_2-(C_1$-C4 alkyl)-;

X: is -CO-;

Y: is -O-, C_1-C_4 alkylene, C_1-C_4 alkynylene, cycloalkylene, aminocarbonyl, -NH-, $-N(C_1$-C_4 alkyl)-, - C_1-C_4 alkylene-$N(C_1$-C_4 alkyl)-, $-CH_2O-$, -CH(OH)- or $-OCH_2-$;

Z: is hydrogen, halogen, nitro, amino, C_1-C_4 alkyl, C_1-C_4 alkoxy, cyano, trifluoromethyl, hydroxyl or carboxyl;

R^1 and R^2: are hydrogen, or together form a C_1-C_3 bridge; and

n and m: independently are 0-3, wherein n and m cannot each be 0, and wherein m+n=4;

or an optical antipode, racemate or pharmaceutically-acceptable salt thereof,
wherein the compound of formula (I) is administered in an amount effective for alleviating at least one symptom of the disease or disorder, and

wherein the pain is caused by a disease or disorder selected from the group consisting of a traumatic injury of a brain or spinal cord, human immunodeficleney virus related neuronal injury, amyotrophic lateral sclerosis, tolerance or dependence to opioid pain treatment, withdrawal syndromes from alcohol, opioids or cocaine, ischemic CNS disorders, chronic neurodegenerative disorders, Alzheimer's disease, Parkinson's disease, Huntington's disease, epilepsy, anxiety, depression, migraine, psychosis, muscular spasm, dementia, hypoglycemia, degenerative disorders of the retina, glaucoma, asthma, tinnitus or hearing loss.